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In re Application of:

LI et al.

Serial No. NOT YET ASSIGNED

Filed: November 17, 2003

For: **IONTOPHORETIC DELIVERY OF ROTIGOTINE FOR THE TRAETMENT OF
PARKINSON'S DISEASE**

REQUEST FOR PRIORITY UNDER 35 U.S.C. §119

Commissioner of Patents
P.O. Box 1450
Alexandria, VA 22313-1450


Commissioner:

In the matter of the above-captioned application, notice is hereby given that the Applicant claims as priority date December 2, 2002, the filing date of the corresponding application filed in European Patent Office, bearing Application Number 02026871.0.

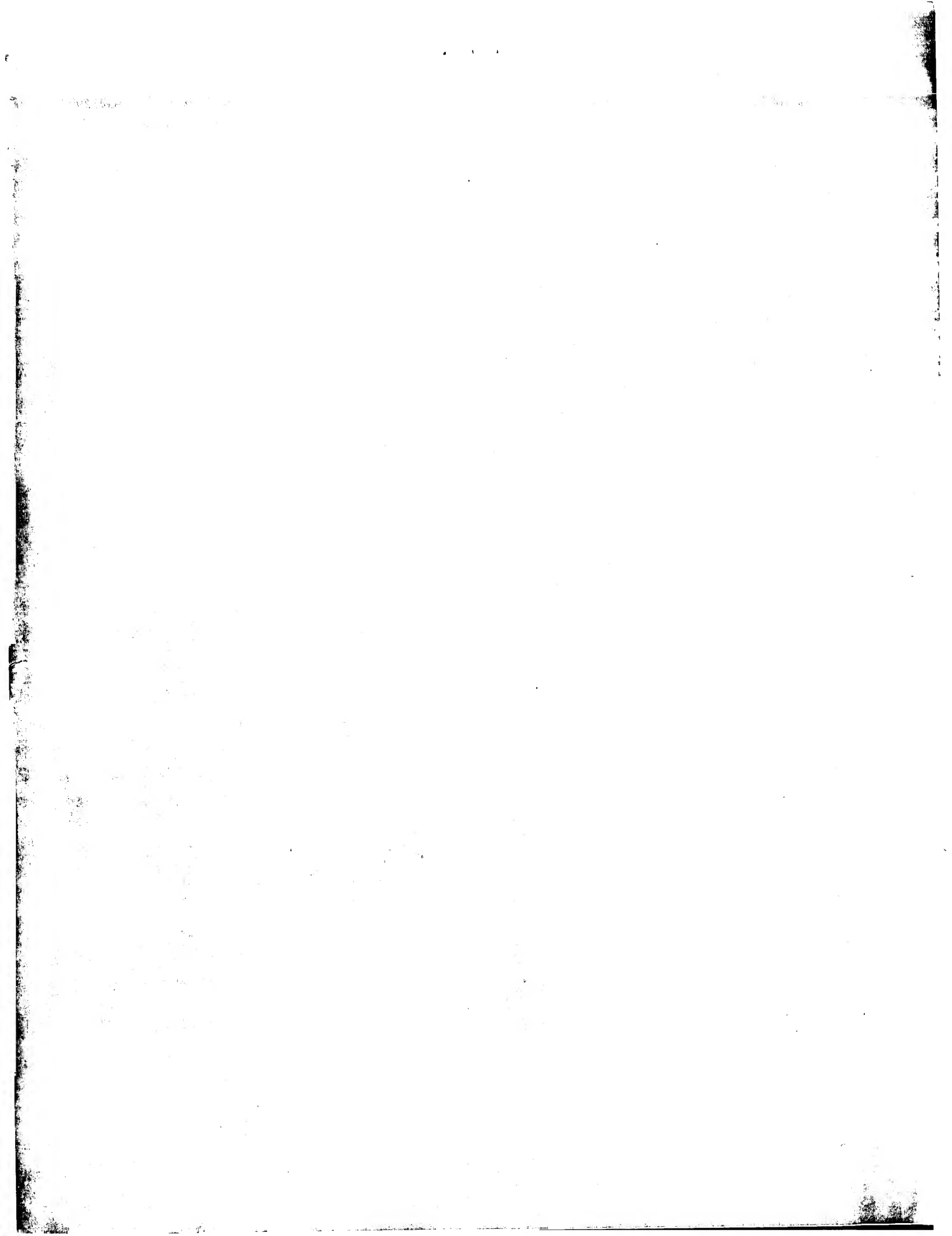
A Certified Copy of the corresponding application is submitted herewith.

Respectfully submitted,
NATH & ASSOCIATES PLLC

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Patentanmeldung Nr. Patent application No. Demande de brevet n°

02026871.0

Der Präsident des Europäischen Patentamts;
Im Auftrag

For the President of the European Patent Office

Le Président de l'Office européen des brevets
p.o.

R C van Dijk



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Bezeichnung der Erfindung/Title of the invention/Titre de l'invention:
(Falls die Bezeichnung der Erfindung nicht angegeben ist, siehe Beschreibung.
If no title is shown please refer to the description.
Si aucun titre n'est indiqué se référer à la description.)

Intophoretic delivery of rotigotine for the treatment of Parkinson's disease

In Anspruch genommene Priorität(en) / Priority(ies) claimed /Priorité(s)
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Parkinson's disease is believed to be primarily caused by the degeneration of dopaminergic neurons in the substantia nigra. This, in effect, results in loss of tonic dopamine secretion and dopamine-related modulation of neuronal activity in the caudate nucleus, and thus in a deficiency of dopamine in certain brain regions. The resulting imbalance of neurotransmitters acetylcholine and dopamine eventually results in disease related symptoms. Although usually regarded as a motor system disorder, Parkinson's disease is now considered to be a more complex disorder that involves both motor and nonmotor systems. This debilitating disease is characterized by major clinical features including tremor, bradykinesia, rigidity, dyskinesia, gait disturbances, and speech disorders. In some patients, dementia may accompany these symptoms. Involvement of the autonomic nerve system may produce orthostatic hypotension, paroxysmal flushing, problems with thermal regulation, constipation, and loss of bladder and sphincter control. Psychological disorders such as loss of motivation and depression may also accompany Parkinson's disease.

Parkinson's disease is primarily a disease of middle age and beyond, and it affects both men and women equally. The highest rate of occurrence of Parkinson's disease is in the age group over 70 years old, where Parkinson's disease exists in 1.5 to 2.5% of that population. The mean age at onset is between 58 and 62 years of age, and most patients develop Parkinson's disease between the ages of 50 and 79. There are approximately 800,000 people in the United States alone with Parkinson's disease.

Early motor deficits of Parkinson's disease can be traced to incipient degeneration of nigral dopamine-releasing cells. This neuronal degeneration produces a defect in the dopaminergic pathway that connects the substantia nigra to the striatum. As the disease progresses, refractory motor, autonomic, and mental abnormalities may develop, which implies that there is progressive degeneration of striatal receptor mechanisms.

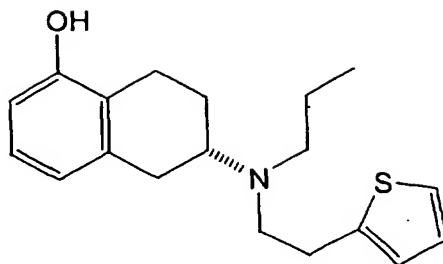
The clinical diagnosis of Parkinson's disease is based on the presence of characteristic physical signs. The disease is known to be gradual in onset, slowly progressive, and variable in clinical manifestation. Evidence suggests that the striatal dopamine content declines to 20% below levels found in age-matched controls before symptoms occur.

Treatment of Parkinson's disease has been attempted with, inter alia, L-dopa (levodopa), which still is the gold standard for the therapy of Parkinson's disease. Levodopa passes the blood-brain barrier as a precursor for dopamine and is then converted into dopamine in the brain. L-dopa improves the symptoms of Parkinson's disease but may cause severe side effects. Moreover, the drug tends to lose its effectiveness after the first two to three years of treatment. After five to six years, only 25% to 50% of patients maintain improvement.

Furthermore a major drawback of currently utilized therapies for Parkinson's disease is the eventual manifestation of the "fluctuation syndrome", resulting in "all-or-none" conditions characterized by alternating "on" periods of mobility with dyskinesias and "off" periods with hypokinesia or akinesia. Patients who display unpredictable or erratic "on-off" phenomena with oral anti-Parkinson therapy have a predictable beneficial response to i.v. administration of L-dopa and other dopamine agonists, suggesting that fluctuations in plasma concentrations of drug are responsible for the "on-off" phenomena. The frequency of "on-off" fluctuations has also been improved by continuous infusions of the dopamine receptor agonists apomorphine and lisuride. However, this mode of administration is inconvenient. Therefore, other modes of administration providing a more constant plasma level, such as topical administration, are beneficial and have been suggested in the past.

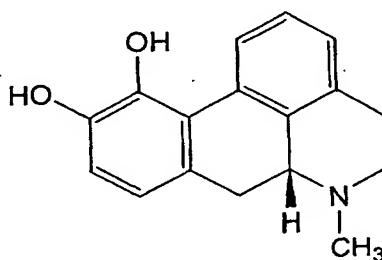
As mentioned above, one treatment approach for Parkinson's disease involves dopamine receptor agonists. Dopamine receptor agonists (sometimes also referred to as dopamine agonists) are substances which, while structurally different from dopamine, bind to different subtypes of dopamine receptors and trigger an effect which is comparable to that of dopamine. Due to the reduced side-effects, it is advantageous when the substances selectively bind to a subgroup of dopamine receptors, i.e. the D2 receptors.

One dopamine receptor agonist which has been used to treat the symptoms of Parkinson's disease is rotigotine. It has mostly been tested in the form of its hydrochloride. Rotigotine is the International Non-Proprietary Name (INN) of the compound (-)-5,6,7,8-tetrahydro-6-[propyl-[2-(2-thienyl)ethyl]-amino]-1-naphthalenol having the structure shown below



It has before been known to administrate rotigotine by passive transdermal therapeutic systems (TTS). Such passive transdermal therapeutic systems for the administration of rotigotine have been described for example in WO 94/07568 and WO 99/49852. However, the rotigotine flux obtained with these passive transdermal therapeutic systems is not necessarily sufficient for all patients.

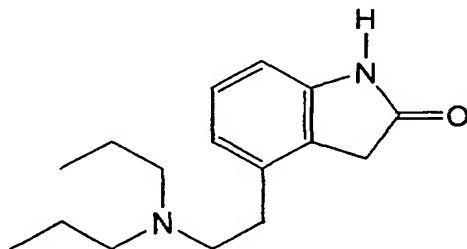
Another dopamine agonist which has been used in the treatment of Parkinson's disease is R-apomorphine. R-apomorphine is the International Non-Proprietary Name (INN) of the compound (R)-5,6,6a,7-tetrahydro-6-methyl-4H-dibenzoquinoline-11,12-diol having the structure shown below



Several approaches to develop a system for iontophoretic administration of R-apomorphine have previously been described (see for example R. van der Geest, M. Danhof, H.E. Boddé "Iontophoretic Delivery of Apomorphine: *In Vitro* Optimization and Validation", *Pharm. Res.* (1997), 14, 1797-1802; M. Danhof, R. van der Geest, T. van Laar, H.E. Boddé, "An integrated pharmacokinetic-pharmacodynamic approach to optimization of R-apomorphine delivery in Parkinson's disease", *Advanced Drug Delivery Reviews* (1998), 33, 253-

263). However, in spite of these efforts, only concentrations at the lower end of the therapeutic concentration range of 1.4 to 10.7 ng/ml could be obtained.

- 5 A further dopamine antagonist is ropinirole hydrochloride. Ropinirole (INN) is (4-[2-dipropylamino)ethyl]-1,3-dihydro-2H-indol-2-one) having the structure shown below



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Although the iontophoretic administration of ropinirole was considered feasible, it was only possible to obtain fluxes at the lower end of the therapeutic range (see A. Luzardo-Alvarez, M. B. Delgado-Charro, J. Blanco-Méndez,
15 "Iontophoretic Delivery of Ropinirole Hydrochloride: Effect of Current Density and Vehicle Formulation", Pharmaceutical Research (2001), 18(12), 1714-1720).

Many patients need concentrations that are significantly
20 higher than the ones feasible using iontophoretic delivery of apomorphine or ropinirole.

In view of the broad range of symptoms of Parkinson's disease and the differing severity, there is a strong demand for a
25 method which allows adjusting the ropinirole flux across the skin and at the same time allows for a constant receptor stimulation of the dopamine receptors of Parkinson patients. Preferably such a system should also allow for ropinirole fluxes higher than the ones achieved by passive transdermal
30 delivery systems.

In view of the discouraging experiences with the iontophoretic delivery of apomorphine, it has been surprising that iontophoretic delivery of rotigotine could provide plasma levels of rotigotine which are not only higher than the ones of conventional passive diffusion systems but are actually in a range that allows for the delivery of pharmaceutically effective drug dosages. The results obtained by using this invention allow for a reasonable expectation that an effective treatment of Parkinson's disease can be provided. It should be understood that the term "treatment" in the context of this application is meant to designate a treatment or alleviation of the symptoms of Parkinson's disease, rather than a real causative treatment leading to a complete cure.

Summary of the Invention

The present invention provides the use of a composition comprising rotigotine and at least one chloride salt in a concentration of 1 to 140 mmol/l, the composition having a pH of 4 to 6.5 for the preparation of a iontophoretic device for the treatment of Parkinson's disease.

Iontophoresis is the introduction of various ions into the skin by means of electricity. If compared to passive transdermal delivery, iontophoresis provides for several advantages which are useful in the treatment of Parkinson's disease:

- it allows programming of the flux at the required therapeutic rate by adjusting the electric current, and
- it permits a rapid start or termination of administration of the medication, if needed, by simply turning the iontophoretic delivery system on or off.

As the iontophoretic flux is influenced by several parameters, it is crucial for achieving an optimal flux to separately optimise these parameters.

5 Surprisingly, it was found that using a composition having a pH of 4 to 6.5 and comprising rotigotine and at least one chloride salt in a concentration of 1 to 140 mmol/l in the donor chamber of the iontophoretic device, fluxes well within the therapeutic range could be achieved.

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By reducing the electrolyte concentration in the donor compartment it was possible to achieve the target iontophoretic flux at lower current density or to increase the transdermal dose per area unit.

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During the studies conducted to evaluate the feasibility of iontophoretic delivery of rotigotine, it was found that the solubility of rotigotine decreases when the pH is increased. However, surprisingly it was found that a therapeutically
20 relevant rate was achieved within the pH interval of 4 to 6.5 at very low rotigotine concentrations.

To provide for an optimal flux across human stratum corneum it was also necessary to provide for a sufficient
25 concentration of Cl^- ions for the electrode reaction in the donor phase. However, while maintaining the electrode reaction the addition of chloride salts reduces the solubility of rotigotine. Thus, a concentration of chloride salts of 1 to 140 mmol/l, preferably 50 to 100 mmol/l, more
30 preferably 60 to 80 mmol/l, proved optimal.

The rotigotine concentration may be varied in accordance with the patient's needs and the flux required for obtaining a therapeutic effect in the treatment Parkinson's disease.

35 However, for a optimal performance it is preferably at least 0.5 mg/ml, more preferably 0.5 mg/ml to 3 mg/ml.

All chloride salts which are pharmaceutically acceptable may be employed in the composition of the invention. In a preferred embodiment of the invention the chloride salt is selected from NaCl, triethylammonium chloride and tributylammonium chloride. Triethylammonium chloride and tributylammonium chloride are especially preferred, because they result in higher fluxes of rotigotine.

In an especially preferred embodiment of the invention the composition, which is used as the donor phase of the iontophoretic device, comprises rotigotine in a concentration of 0.5 to 3 mg/ml and at least one of triethylammonium chloride and tributylammonium chloride in a concentration of 60 to 80 mmol/l, the donor phase has a pH of 4.5 to 5.5.

In another aspect the present invention provides a method for the treatment of Parkinson's disease, wherein a iontophoretic device, which comprises a composition comprising rotigotine and at least one chloride salt in a concentration of 1 to 140 mmol/l, the composition having a pH of 4 to 6.5, is applied onto the skin of a patient in need thereof.

Any conventional iontophoretic device may be used in the invention. Such iontophoretic devices are described e.g. in V. Nair, O. Pillai, R. Poduri, R. Panchagnula, "Transdermal Iontophoresis. Part I: Basic Principles and Considerations" Methods Find. Exp. Clin. Pharmacol. (1999), 21(2), 139-151.

The current density employed during iontophoresis may be varied according to the patient's needs and will depend on the iontophoretic device and the composition used. A suitable current may be determined by the attendant physician. In general, a suitable current density will be in the range of preferably 200 to 500 $\mu\text{A}/\text{cm}^2$.

Example 1

In vitro iontophoretic studies for the administration of rotigotine were carried out with three-chamber flow-through diffusion cells as described by R. van der Geest et al. (R. van der Geest, M. Danhof, H.E. Boddé, "Validation and testing of a new iontophoretic continuous flow through transport cell", J. Control. Release (1998), 51, 85-19). On both sides of the acceptor compartment human stratum corneum (SC) was situated. A dialysis membrane having a 5.000 Da cut-off was used as supporting membrane. The volume of the outer chambers was approximately 2 ml, while the volume of the acceptor compartment was 0.54 ml. The two outer chambers contained the silver plate (anode) or silver/silver chloride (cathode) driver electrodes. The donor phase consisted of rotigotine solution buffered with 5 mM citrate buffer (2.1 mM sodium citrate dihydrate and 2.9 mM citric acid).

Using this set-up, a pH in the donor chamber of 5, a current density of 500 $\mu\text{A}/\text{cm}^2$, a pH in the acceptor chamber of 7.4, a temperature of 20°C and an NaCl concentration in the donor chamber of 70 mmol/l, the flux of rotigotine was measured for different drug concentrations in the donor phase.

Rotigotine conc. (mg/ml) (Donor solution)	Rotigotine conc. (mM) (Donor solution)	Flux (nmol/cm ² /h) Rotigotine
0.5	1.4	22.9
1.0	2.8	30.2
1.4	3.98	53.2

Example 2:

Using a similar procedure as in Example 1 and a concentration of rotigotine of 1.4 mg/ml (3.98 mM), a pH in the donor

chamber of 5, a current density of $500 \mu\text{A}/\text{cm}^2$, a pH in the acceptor chamber of 7.4 and a temperature of 20°C , but substituting triethylammonium chloride (TEACl) or tributylammonium chloride (TBACl) for NaCl, the influence of the different cations on the flux was evaluated. The concentration of the chloride salts in the donor solution was $70 \text{ mmol}/\text{l}$.

Co-ions source	Flux ($\text{nmol}/\text{cm}^2/\text{h}$) Rotigotine
NaCl	53.2
TEACl	72.8
TBACl	62.0

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Example 3:

Using a similar procedure and the same parameters as in Example 2, the influence of reducing the pH in the acceptor chamber from 7.4 to 6.2 was evaluated for different chloride salts. The concentration of the chloride salts in the donor solution was $70 \text{ mmol}/\text{l}$.

Co-ions source	Flux ($\text{nmol}/\text{cm}^2/\text{h}$) Rotigotine
NaCl	58.9
TEACl	43.2
TBACl	76.5

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Claims

1. Use of a composition comprising rotigotine and at least one chloride salt in a concentration of 1 to 140 mmol/l, the composition having a pH of 4 to 6.5 for the preparation of a iontophoretic device for the treatment of Parkinson's disease.
2. Use according to claim 1, wherein the concentration of rotigotine is at least 0.5 mg/ml.
3. Use according to any one of claims 1 or 2, wherein the concentration of rotigotine is 0.5 to 3 mg/ml.
4. Use according to any one of claims 1 to 3, wherein the chloride salt is selected from NaCl, triethylammonium chloride and tributylammonium chloride.
5. Use according to claim 4, wherein the chloride salt is triethylammonium chloride or tributylammonium chloride.
6. Use according to any one of claims 1 to 5, wherein the concentration of the chloride salt is 60 to 80 mmol/l.
7. Use according to any one of claims 1 to 6, wherein the composition is used in the donor phase of the iontophoretic device.
8. Use according to any one of claims 1 to 7, wherein the composition in the donor phase of the iontophoretic device comprises rotigotine in a concentration of 0.5 to 3 mg/ml and at least one of triethylammonium chloride and tributylammonium chloride in a concentration of 60 to 80 mmol/l, wherein the pH of the donor phase is 4.5 to 5.5.

9. A method for the treatment of Parkinson's disease characterised by applying a iontophoretic device, which comprises a composition comprising rotigotine and at least one chloride salt in a concentration of 1 to 140 mmol/l, the composition having a pH of 4 to 6.5, onto the skin of a patient in need thereof.

EPO - Munich
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Abstract

By using a composition comprising rotigotine and at least one chloride salt in a concentration of 1 to 140 mmol/l, the
5 composition having a pH of 4 to 6.5 in a iontophoretic device for the treatment of Parkinson's disease, it became possible to obtain a rotigotine flux across the human stratum corneum which was higher than the one previously obtained with conventional passive diffusion systems.

